## 09/902556

FILE 'REGISTRY' ENTERED AT 10:54:29 ON 31 JAN 2003 L1 1 S GSSFLSPEAKLOPR/SQSP ANSWER 1 OF 1 REGISTRY COPYRIGHT 2003 ACS L1392687-99-9 REGISTRY RN 6 SSFLSPE L-Arginine, glycyl-L-seryl-O-(1-oxooctyl)-L-seryl-L-phenylalanyl-L-CN leucyl-L-seryl-L-prolyl-L-.alpha.-glutamyl-L-alanyl-L-lysyl-L-leucyl-L-glutaminyl-L-prolyl- (9CI) (CA INDEX NAME) OTHER NAMES: 3: PN: WO0208250 SEQID: 3 claimed protein EP 00774 CN 14 SQL SEQ 1 GSSFLSPEAK LQPR HITS AT: 1-14 REFERENCE 1: 137:73541 REFERENCE 2: 136:129068 FILE 'HCAPLUS' ENTERED AT 10:55:18 ON 31 JAN 2003 L2 2 S L1 L2 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2003 ACS 2002:317047 HCAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 137:73541 Short ghrelin peptides neither displace ghrelin TITLE: binding in vitro nor stimulate GH release in vivo AUTHOR(S): Torsello, Antonio; Ghe, Corrado; Bresciani, Elena; Catapano, Filomena; Ghigo, Ezio; Deghenghi, Romano; Locatelli, Vittorio; Muccioli, Giampiero 400 nur Department of Experimental and Environmental CORPORATE SOURCE: Medicine and Biotechnologies, University of Milano-Bicocca, Milan, Italy Endocrinology (2002), 143(5), 1968-1971 SOURCE: CODEN: ENDOAO; ISSN: 0013-7227 Endocrine Society PUBLISHER: DOCUMENT TYPE: Journal LANGUAGE: English Ghrelin is an acylated peptide recently isolated from rat stomach AB that potently stimulates GH release in vitro and in vivo in rat and Ghrelin specifically activates the receptor for the growth hormone secretagogues (GHS-Rla), and it has been proposed as the endogenous ligand mimicked by these synthetic compds. Very recently, it was shown in cells transfected with the GHS-Rla that short acylated peptides encompassing the first 4-5 residues of ghrelin were capable of increasing intracellular calcium almost as efficiently as the full-length ghrelin. In the present study, we demonstrate that truncated analogs of ghrelin are ineffective in stimulating GH release in neonatal rats and do not displace radiolabeled ghrelin from binding sites in membranes from human hypothalamus and pituitary. In conclusion, our data demonstrate

Searcher: Shears 308-4994

that the ability of short ghrelins to stimulate the GHS-Rla in

GH secretion in vivo.

transfected cells is not predictive of their capability to stimulate

## 09/902556

392687-99-9, EP 00774 TΤ

> RL: BSU (Biological study, unclassified); BIOL (Biological study) (ghrelin short peptides neither displace ghrelin binding in vitro

nor stimulate growth hormone release in vivo)

23

REFERENCE COUNT:

THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE

IN THE RE FORMAT

ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER:

2002:90068 HCAPLUS

DOCUMENT NUMBER:

136:129068

TITLE:

Ghrelin antagonist peptides

INVENTOR(S):

Deghenghi, Romano

PATENT ASSIGNEE(S):

Zentaris A.-G., Germany

SOURCE:

PCT Int. Appl., 9 pp. CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.				KIND		DATE		APPLICATION NO.					DATE				
	WO 2002008250 WO 2002008250								WO 2001-EP7929 2001071							0710	
	V	v: .	AU,	BG,	BR,	BY,	CA,	CN,	co,	CZ,	EE,	GE,	HR,	HU,	ID,	IL,	IN
			IS,	JP,	KG,	KR,	KZ,	LT,	LV,	MK,	MX,	NO,	NZ,	PL,	RO,	RU,	SG
			SI,	SK,	TR,	UA,	UZ,	YU,	ZA,	AM,	AZ,	BY,	KG,	ΚZ,	MD,	RU,	TJ
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OMURD GOUDGE (G). MADDAM 126-120060																	

OTHER SOURCE(S): MARPAT 136:129068

Novel peptides are disclosed having antagonistic properties to the Growth Hormone releasing peptide known as Ghrelin. The new peptides are useful in decreasing the circulating levels of Growth Hormone in a mammal and have therapeutic value. Peptide Gly-Ser-Ser(Octanoyl)-Phe, prepd. by solid phase synthesis, antagonized the effect of ghrelin by reducing growth hormone release in 10-day old rats.

392687-99-9P

RL: PAC (Pharmacological activity); PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(ghrelin antagonist peptides)

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